Whole-proteome genetic analysis of dependencies in assembly of a vertebrate kinetochore

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Kinetochores orchestrate mitotic chromosome segregation. Here, we use quantitative mass spectrometry of mitotic chromosomes isolated from a comprehensive set of chicken DT40 mutants to examine the dependencies of 93 confirmed and putative kinetochore proteins for stable association with chromosomes. Clustering and network analysis reveal both known and unexpected aspects of coordinated behavior for members of kinetochore protein complexes. Surprisingly, CENP-T depends on CENP-N for chromosome localization. The Ndc80 complex exhibits robust correlations with all other complexes in a “core” kinetochore network. Ndc80 associated with CENP-T interacts with a cohort of Rod, zw10, and zwilch (RZZ)-interacting proteins that includes Spindly, Mad1, and CENP-E. This complex may coordinate microtubule binding with checkpoint signaling. Ndc80 associated with CENP-C forms the KMN (Kn1, Mis12, Ndc80) network and may be the microtubule-binding “workhorse” of the kinetochore. Our data also suggest that CENP-O and CENP-R may regulate the size of the inner kinetochore without influencing the assembly of the outer kinetochore.

Introduction

Genetic and biochemical analyses have led to identification of ~100 proteins that localize at centromeres and kinetochores (Takahashi et al., 1994; Obuse et al., 2004; Foltz et al., 2006; Okada et al., 2006; Ohta et al., 2011; Perpelescu and Fukagawa, 2011; Biggins, 2013; Tanaka, 2013). A group of 16 proteins known as the CCAN (constitutive centromere-associated network) comprises the inner, chromatin-proximal portion of the kinetochore (Cheeseman and Desai, 2008; Westhorpe and Straight, 2013; Fukagawa and Earnshaw, 2014). CCAN components remain associated with centromeres across the cell cycle and are a foundation for outer kinetochore assembly during mitosis. The CCAN is composed of discrete subcomplexes (De Wulf et al., 2003; Foltz et al., 2006; Okada et al., 2006; Hori et al., 2008b; Nishino et al., 2012). The precise interactions among CCAN proteins and complexes in chromosomes are being studied in many laboratories, and several models have been proposed (DeLuca and Musacchio, 2012; Fuchinetti et al., 2013; Cheeseman, 2014; Fukagawa and Earnshaw, 2014; Westhorpe and Straight, 2015; Kim and Yu, 2015).

The mid and outer kinetochore are also characterized by the presence of protein complexes, including CENP-O/-P/-Q/-R/-U (throughout, we will use this shorthand when listing multiple CENP proteins, e.g., CENP-O, CENP-P, and CENP-Q), Ndc80, Mis12, Rod, zw10, and zwilch (RZZ), and the Nup107/Nup160 nucleoporin complexes. The Ndc80 and Mis12 complexes, together with the protein KNL-1, form the KMN network, which is a major site of microtubule binding in the outer kinetochore (Cheeseman et al., 2006).

Some of these complexes have been isolated and characterized structurally. In addition to several CENP-A nucleosome-related structures (Sekulic et al., 2010; Tachiwana et al., 2011; Kato et al., 2013), structures have also been reported for the dimerization domain of CENP-C (Cohen et al., 2008), a bonsai form of the Ncd80 complex (Ciferri et al., 2008), as well as several of the protein domains involved in that complex (Wei et al., 2006, 2007), the CENP-T-W-S-X complex (Nishino et al., 2012), the Knl1/Mis12 complex (Petrovic et al., 2014), the lml3/Chl4 heterodimer (Hinshaw and Harrison, 2013), and the Ska complex (Jeyaprakash et al., 2012). However, functional analysis of these complexes is complicated, because in chromosomes, they form an insoluble network. Thus, studies of the functions and dependencies of the proteins in their “native environment” (e.g., in situ in mitotic chromosomes) are crucial.

Here, we combined quantitative proteomics using stable isotope labeling by amino acids (SILAC) technology (Ong et
al., 2002) with DT40 genetics to map a vertebrate kinetochore by monitoring the behavior of the entire mitotic chromosome proteome after the depletion of defined components. We previously showed that this approach can track the behavior of individual protein complexes in the presence of hundreds or thousands of other proteins (Ohta et al., 2010).

Our studies define dependency pathways for the stable association of a number of protein complexes with mitotic chromosomes. Several unexpected relationships were found. The data suggest that the organization of kinetochore proteins into functional complexes may be more complicated than suggested by previous studies.

Results

Strategy and overview

Analysis of the proteome of DT40 mitotic chromosomes by mass spectrometry (Ohta et al., 2010) enabled us to both make a list of proteins that associate with mitotic chromosomes and also unearth specific protein–protein interactions within and between protein complexes (Ohta et al., 2010; Samejima et al., 2012). Here, we used similar unbiased whole-proteome quantification to define dependency relationships for kinetochore protein assembly.

Several caveats should be kept in mind when comparing our data with other studies. First, because chromosome isolation requires a very high mitotic index (>80%), cultures must be healthy enough to continue to cycle. Therefore, most of our experiments with conditional knockout (KO) cells (with the exception of null alleles of nonessential genes) were performed before the target protein was completely lost. Second, in order to obtain a mitotic index of >80%, cultures were blocked for 12 h with nocodazole. This may induce some alterations in the kinetochore, but all of our experiments compare wild-type and mutant chromosomes subjected to an identical treatment. Third, our analysis detected changes in the entire proteome of mitotic chromosomes, not only at kinetochores. This is potentially significant, as despite the apparent specificity of antibody staining, a recent study reported that ~70% of CENP-A is located at noncentromeric sites in RPE-1 cells (Bodor et al., 2014). The ratio of centromeric to noncentromeric binding is not known for other kinetochore proteins.

We isolated mitotic chromosomes from 11 DT40 mutant cell lines for whole-proteome analysis. Eight of those lines are conditional knockouts where the target gene is disrupted and cell lines for whole-proteome analysis. Eight of those lines are essential genes encoding CENP-O, CENP-R, or CENP-S were deleted (referred to as CENP- OME; Ong et al., 2002). The relative abundance of proteins in the same clusters (Fig. 1 and Table S1). This included components of the Ndc80, Mis12, RZZ, chromosome passenger complex (CPC), anaphase-promoting complex/cyclosome, Nup107/Nup160 nucleoporin, dynein, and dynactin complexes (Fig. 1). Meta-analysis of variations in the kinetochore proteome performed by combining results from the different genetic perturbations also allowed us to identify novel cohorts of kinetochore proteins. We define a “cohort” as a group of proteins that exhibit coordinated behavior in response to diverse genetic perturbations but have not been independently shown to form a protein complex.

Hierarchical clustering analysis of the H/L ratios across all experiments identified two clusters of proteins that appeared to correspond to contaminants and hitchhikers (red lines at left in Fig. 1). Hitchhikers were defined in Ohta et al. (2010) as cytoplasmic proteins that bind to mitotic chromosomes after nuclear envelope breakdown but do not have a role in mitotic chromosome structure or function. These and other proteins whose variation correlated with the tissue culture medium in which cells were grown were excluded from our statistical analysis as explained in Materials and methods.

A subset of CCAN proteins is depleted from CENP-1OFF and CENP-HOFF chromosomes

As a first approach to determining the effects of disrupting the CCAN on kinetochore composition, we determined the proteome of chromosomes isolated from CENP-1OFF or CENP-HOFF cells. CENP-I and CENP-H are closely interacting CCAN members (Nishihashi et al., 2002; Okada et al., 2006; Basilico et al., 2014).

In a typical experiment with CENP-1OFF chromosomes, 1,504 proteins were quantified. Depletion of CENP-I was efficient, with ~12% of the wild-type level remaining on mutant chromosomes (H/L ratio = 8.09). CENP-H/K-L/-N/-T were also significantly depleted, with <16% of wild-type levels remaining (Fig. 2, A and B; and Fig. S1 D). In contrast, CENP-A and CENP-C were little affected, with >72% remaining on the chromosomes.

A scatterplot of the H/L ratios of 1,421 proteins from two replicate experiments shows that these observations were reproducible (Fig. 2 B). In these plots of H/L ratios in experiment i versus H/L ratios in experiment ii, good correlation is shown when points cluster along a tight diagonal line. This plot reveals that CENP-H/K-L/-N/T are the proteins most affected by CENP-I depletion, with H/L ratios nearly as large as CENP-I (Fig. 2 B). These proteins form a discrete cluster separate from the rest of the population, suggesting that they comprise one or more protein complexes that are not stably associated with CENP-I OFF chromosomes. CENP-S/-X were also significantly depleted from these chromosomes, and they formed a distinct cluster.
CENP-O/-P/-Q/-R/-U were somewhat less depleted and exhibited a tight cluster of H/L ratios (Fig. 2 B, inset). Outer kinetochore proteins, including members of the Ndc80 complex, clustered with similar H/L ratios, and members of the RZZ and Mis12 complexes were also among the top 5% most reduced proteins (above and to the right of the two orthogonal blue lines).

Similar results were obtained after CENP-H depletion. Fig. 2 C shows a scatterplot of the H/L ratio for 1,515 proteins from two replicate experiments with CENP-H\textsuperscript{OFF} chromosomes.

In summary, depletion of CENP-I or CENP-H from mitotic chromosomes caused a reproducible downstream pattern of depletion of both inner and outer kinetochore proteins from isolated mitotic chromosomes.

Similar proteomes were observed with chromosomes from CENP-H\textsuperscript{OFF}, CENP-I\textsuperscript{OFF}, and CENP-K\textsuperscript{OFF}, or CENP-M\textsuperscript{OFF} cells. These experiments involved numerous repetitions of proteomics experiments conducted with multiple mutants. A correlation matrix was plotted to compare the results of different experiments (Fig. S2, A and B). The Pearson correlation coefficient was calculated from the distribution of all quantified proteins on chromosomes (gray), all centromere proteins (blue), or CCAN proteins only (red; Fig. S2, C–H). A strong correlation was observed between replicate experiments with the same cell line and also between the CENP-H and CENP-I depletion results (Figs. 3 A and S2), confirming that both mutants have a similar
Figure 2. Behavior of CENP-H,-I,-K,-M,-N,-L,-T in CENP-I and CENP-H KOs. (A) Histograms show the distribution of log$_2$(H/L ratio) of all proteins quantified, ribosomal proteins, and centromere proteins. CCAN proteins (red) are shown as stacked columns. Proteins with a positive score (rightwards on the abscissa) were depleted from CENP-I OFF chromosomes. A blue line designates the top 5%. Those with a negative score (leftward on the abscissa) are accumulated in CENP-I OFF chromosomes. (B) CENP-I OFF: H/L ratios of centromere proteins (blue), CCAN proteins (red), and others (gray) from two replicate experiments are plotted against one another on a log scale in this scatterplot. The blue lines designate the top 5% most depleted proteins. Proteins that decreased in the CENP-I OFF chromosomes map to the top right, and those that increased map to the bottom left. Enlarged plot of the areas boxed in left panel is shown in the blue box on the right. CCAN proteins (red) are designated with the prefix “CENP-” removed. Members of the Ndc80 complex are in blue, whereas members of the Mis12 complex and KNL1 are in green. (C) CENP-H OFF: H/L ratios from two replicate experiments are plotted against one another. Abbreviations and color-coding are as in B. A magnified plot of the boxed area is shown in a blue box. Note that to appear in scatterplots here and below, proteins must have been identified in both experiments being compared. Very small proteins that give rise to few peptides, such as CENP-M, CENP-S, and CENP-X, were occasionally not detected and are therefore not shown in some of the plots. wt, wild type.
effect on the behavior of centromere proteins, in particular CCAN proteins (Figs. 3 C and S2).

A similar correlation was obtained for comparisons involving CENP-K\textsuperscript{OFF} and CENP-M\textsuperscript{OFF}. In all, proteomes of chromosomes from CENP-H\textsuperscript{OFF}/CENP-K\textsuperscript{OFF}/CENP-M\textsuperscript{OFF} cells form one closely correlated group (top left yellow box in Fig. 3 C), consistent with them comprising a protein complex (Basilico et al., 2014). Similarly, proteomes of chromosomes from CENP-O\textsuperscript{KO} and CENP-R\textsuperscript{KO} form a distinct group (Fig. 3 C, bottom right yellow box).

Importantly, these correlations were diminished when cross-comparisons were made between proteomes of chromosomes depleted of proteins associated with different kinetochore subcomplexes (Figs. 3 B and S2). For example, proteome comparisons between chromosomes isolated from CENP-C\textsuperscript{OFF}, CENP-C\textsuperscript{S\textsuperscript{KO}}, CENP-H\textsuperscript{OFF}/CENP-I\textsuperscript{OFF}/CENP-K\textsuperscript{OFF}/CENP-M\textsuperscript{OFF}/CENP-N\textsuperscript{OFF}, and CENP-O\textsuperscript{KO}/CENP-R\textsuperscript{KO} cells showed a significantly reduced correlation. This indicates that these four groups of proteins affect the overall chromosomal association of centromere proteins in different ways.

CENP-T behaves coordinately with CENP-N and CENP-L

CENP-N was originally identified as a CCAN member (Foltz et al., 2006; Izuta et al., 2006; Okada et al., 2006) that associates with CENP-A nucleosomes (Carroll et al., 2009; Hellwig et al., 2011). CENP-N, CENP-L, and CENP-T were by far the most depleted proteins in CENP-N\textsuperscript{OFF} chromosomes (Fig. 4 A). CENP-H and CENP-I were also significantly decreased but had H/L ratios in a separate (less depleted) cluster together with members of the CENP-O/-P/-Q/-U complex, the Ndc80 complex, the RZZ complex, Mad1, and CENP-E (Fig. 4 A, inset).

These data suggest that CENP-N might be part of a protein cohort with CENP-L and CENP-T that interacts with CENP-H/-I/-K/-M to form a larger protein assembly in intact mitotic chromosomes. To reiterate, we use the term cohort to describe a group of proteins whose correlated behavior is consistent with them forming a protein complex that has yet to be independently confirmed.

CENP-H/-I/-K/-M and CENP-N/-L had very similar profiles in virtually all experiments, with the exception of CENP-N\textsuperscript{OFF} cells (Fig. 4 B). (A “profile plot” is an illustrative summary of changes in the abundance of proteins across all experiments [see Materials and methods]. The gray lines plot levels of all 1,409 proteins analyzed in all experiments.) An unexpected result was obtained when comparing the CENP-N\textsuperscript{OFF} proteome against its CENP-H\textsuperscript{OFF} or CENP-P\textsuperscript{OFF} counterparts. Despite CENP-N being part of the CENP-H/-I/-K/-M/-N cluster, when proteomes comprising only CCAN proteins were compared, those derived from CENP-N\textsuperscript{OFF} chromosomes did not exhibit strong correlations with the others. This is seen as scatter away from the diagonal in Fig. 3 B and a lower correlation in the matrix of Fig. 3 C. This suggests that CENP-N is not always associated with CENP-H/-I/-K/-M.

The profile plots for CENP-N and CENP-L are nearly identical, consistent with studies that these proteins form a heterodimeric complex when they are loaded on chromosomes (Carroll et al., 2009; Hinshaw and Harrison, 2013). Both proteins were significantly depleted from chromosomes isolated from CENP-H\textsuperscript{OFF}/CENP-I\textsuperscript{OFF}/CENP-K\textsuperscript{OFF}/CENP-M\textsuperscript{OFF}/CENP-N\textsuperscript{OFF} cells. Their reduction was, however, more modest in chromosomes from other CCAN mutant cells, i.e., CENP-C\textsuperscript{OFF}/CENP-C\textsuperscript{S\textsuperscript{KO}}/CENP-R\textsuperscript{KO} (Fig. 4 B).

Unexpectedly, the profile plot for CENP-T was strikingly similar to CENP-N and CENP-L except in CENP-S\textsuperscript{KO} experiments (Fig. 4 B). CENP-T was one of the most depleted proteins in chromosomes from CENP-H\textsuperscript{OFF}/CENP-I\textsuperscript{OFF}/CENP-K\textsuperscript{OFF}/CENP-M\textsuperscript{OFF}/CENP-N\textsuperscript{OFF} cells (Fig. 4, B and C). This suggests a strong dependency of CENP-T on CENP-H/-I/-K/-M/-N, particularly on CENP-N and CENP-L. Although CENP-T on its own has a DNA binding activity in vitro (Hori et al., 2008a), its centromere association may also require CENP-H/-I/-K/-M/-N. Indeed a reduction of CENP-T levels in CENP-H\textsuperscript{OFF} cells was reported previously (Hori et al., 2008a). CENP-T and CENP-W make a DNA binding complex with their histone fold domains (Hori et al., 2008a), and both proteins show essentially identical profiles in these experiments (Fig. 4 C).
**CENP-S and CENP-X may function in part independently of CENP-T**

In addition to a CENP-T/-W dimer, a CENP-T/-W/-S/-X tetramer has also been reported at centromeres (Nishino et al., 2012). The profile plot of CENP-S, however, did not always follow that of CENP-T in isolated mitotic chromosomes (Fig. 4C). For instance, CENP-S was less affected than CENP-T by depletion of CENP-H/-I/-N, suggesting that CENP-S/-X may participate in another complex. Indeed, CENP-S/-X also associates with FANCM (Singh et al., 2010; Yan et al., 2010). For comparison, protein profiles for known complexes, such as the four core histones, the cohesin complex and CPC appear as single lines (Fig. S3).

Because CENP-S is dispensable for growth in DT40 cells, we could obtain a complete KO cell line (Amano et al., 2009). CENP-S peptides were not detected on the chromosomes from CENP-S KO cells. This results in an H/L SILAC ratio of zero or infinite value, and the H/L ratios for individual CENP-S peptides were sometimes dealt with as missing values by the Andromeda search engine (i.e., disregarded in calculating the H/L ratio for a given protein). We therefore reviewed the data manually, and the MS1 spectrum of one representative peptide (AAV HYTTGCLQDVADKVLFSK, charge +4) was identified with heavy-labeled species (i.e., from wild-type cells; Fig. 5A, orange boxes). We observed no trace of the corresponding light-labeled peptide, confirming the absence of CENP-S from CENP-S KO chromosomes.

By far, the most strongly affected protein in CENP-S KO chromosomes was its binding partner, CENP-X (Amano et al., 2009), with <10% remaining (Fig. 5B). Inspection of MS1 spectra of a representative CENP-X peptide (QAQAEDLE KVDIEHVEK, charge +4) confirmed that this peptide was virtually absent (data not shown). After CENP-X, the next most depleted centromere proteins were CENP-T and CENP-W, but their depletion was much less extreme. Thus CENP-S KO chromosomes still retain some functional CENP-T/-W dimer. This may explain the viability of CENP-S KO cells.

CENP-N,-L were not reduced as much as CENP-T/-W on CENP-S KO chromosomes (Figs. 4B and 5B). Rather, the H/L ratio of CENP-N and CENP-L fell in the same cluster as CENP-H/-I/-K/-M and close to CENP-P/-Q/-R/-U (Fig. 5B). This suggests that CENP-N and CENP-L are upstream of CENP-T for localization on chromosomes.

Interestingly, outer kinetochore components, including the KMN, RZZ, and Ska complexes plus CENP-E and Mad1, were more depleted than CCAN components on chromosomes lacking CENP-S (Fig. 5B). Indeed, CENP-T can recruit the Ndc80 complex to chromosomes (Hori et al., 2013; Nishino et al., 2013).

**CENP-C is required for association of the KMN network with chromosomes**

CENP-C links the centromeric chromatin with the outer kinetochore. The C-terminal and central regions of CENP-C interact with CENP-A nucleosomes (Kato et al., 2013), the N terminus binds to Mis12 in the KMN network (Carroll et al., 2010; Przewloka et al., 2011; Screpanti et al., 2011), and the PEST region binds CENP-H/-I/-K/-M (Klare et al., 2015). Profile plots showed only modest changes in CENP-C abundance after depletion of other CCAN proteins (Fig. 6A). Notably, significant...
CENP-C associates with chicken mitotic chromosomes independently of other CCAN proteins. CENP-C depletion caused a reduction in Mis12 complex proteins and KNL1 to an extent similar to that of the Ndc80 complex (Fig. 6 C). They formed a grouping distinct from the cluster of CENP-H/-I/-K/-M/-N/-L/-T on the scatter-plot. Although CENP-H/-I/-K/-M/-N/-L/-T were reduced, their abundance on CENP-COFF chromosomes remained about half of the wild-type level.

CENP-C depletion caused a reduction in Mis12 complex proteins and KNL1 to an extent similar to that of the Ndc80 complex (Fig. 6, B and C). This stoichiometric codepletion is consistent with a direct interaction between CENP-C and the Mis12 complex (Przewloka et al., 2011; Screpanti et al., 2011) and contrasts with depletion of CENP-H/-I/-K/-M/-N/-L/-T, where Ndc80 levels fell with much less effect on the Mis12 complex (Fig. 6 C).

Proteins engaged in a protein complex should show coordinated behavior (i.e., low variance in their SILAC ratios) in our experiments. The variance of the SILAC ratios for the Ndc80 and Mis12 subcomplexes and KNL1 was small in chromosomes from mis18αOFF, CENP-COFF, SROJ-OROJ-RROJ cells (Fig. 6 D). This suggests that components of the KMN network holocomplex behaved together as a single entity in those mutants. In contrast, the variance between the subcomplexes was significantly larger in chromosomes from CENP-H/-I/-K/-M/-N/-L/-T, where Ndc80 levels fell with much less effect on the Mis12 complex (Fig. 6 C).

Reduced CENP-A in HJURP̂OFF or Mis18αOFF chromosomes (see Fig. 7) did not impact on CENP-C levels. We conclude that CENP-C associates with chicken mitotic chromosomes independently of other CCAN proteins.

CENP-C expression was turned off by treatment with doxycycline for 48 h. The H/L ratio for CENP-C reveals that only 14% remained on the chromosomes. No other CCAN proteins were similarly affected. After CENP-C, the most depleted proteins in CENP-COFF chromosomes were members of the KMN network (Fig. 6 B). They formed a grouping distinct from the cluster of CENP-H/-I/-K/-M/-N/-L/-T on the scatter-plot. Although CENP-H/-I/-K/-M/-N/-L/-T were reduced, their abundance on CENP-COFF chromosomes remained about half of the wild-type level.

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CENP-N/-L/-T/-S were the next most depleted centromere proteins after CENP-A, in HJURP\textsuperscript{OFF} chromosomes. KIF2C (MCAK), Mis18\textalpha{}, and Mis18\textbeta{} accumulated on chromosomes from HJURP\textsuperscript{OFF} cells (Fig. 7 A). Mis18BP1/KNL2 was essentially unaffected (Fig. 7 A and C).

Mis18\textalpha{} works together with Mis18\textbeta{} and Mis18BP1 in telophase to prime sites for CENP-A recruitment on chromosomes (Fujita et al., 2007). Mis18\textalpha{} and Mis18\textbeta{} were the most depleted proteins when Mis18\textalpha{} transcription was silenced (Fig. 7 B). Mis18BP1 was also significantly depleted, though much less so than Mis18\textalpha{} or Mis18\textbeta{}. Profile plots of the three proteins imply that Mis18BP1 may not associate with Mis18\textalpha{} and Mis18\textbeta{} in chromosomes isolated from nocodazole-treated cells (Fig. 7 C). After Mis18\textalpha{} and Mis18\textbeta{}, CENP-A was the most depleted centromere protein in mis18\textalpha{}\textsuperscript{OFF} chromosomes (Fig. 7 B). CENP-N/-L/-T/-W also showed a significant decrease, with CENP-H/-I/-K being less affected. Interestingly, chromosomal levels of CENP-C were little affected by the reduction of CENP-A in HJURP\textsuperscript{OFF} or Mis18\textalpha{}\textsuperscript{OFF} chromosomes despite the fact that CENP-C has been reported to interact with CENP-A (Carroll et al., 2010).

Taken together, reduction in the amount of CENP-A coincided well with the reduction of CENP-N/-L/-T/-W. This observation is reminiscent of the results of CENP-N depletion, where CENP-N/-L/-T were the most affected proteins.

Accumulation of CCAN proteins on CENP-O\textsuperscript{KO} chromosomes

CENP-O\textsuperscript{KO} and CENP-R\textsuperscript{KO} cells are viable (Okada et al., 2006). CENP-O forms a complex with CENP-P/-Q/-U and CENP-R (Hori et al., 2008b), and these proteins were significantly depleted from CENP-O\textsuperscript{KO} chromosomes (Fig. 8, A and C). CENP-O/-P/-Q/-U were depleted to a somewhat lesser extent from CENP-R\textsuperscript{KO} chromosomes (H/L ratio = ∼2; Fig. 8, C and D). The tight cluster suggests that CENP-O/-P/-Q/-U were lost as a single entity in the absence of CENP-R (though their depletion...
was less severe than seen with loss of CENP-O). Interestingly, CENP-H/-I/-K/-N/-L/-T/-S accumulated on chromosomes isolated from CENP-O$^{KO}$ and CENP-R$^{KO}$ cells (Fig. 8, B and D). This could reflect an enlargement of the inner kinetochore, as cells adapt to continued growth without CENP-O/-P/-Q/-R/-U, either as a direct effect or possibly by modulating the expression or stability of other proteins. Importantly, this accumulation of CCAN proteins was not accompanied by an increase in KMN and RZZ complex components. Thus, increasing the content of CCAN proteins does not necessarily increase the size of the outer kinetochore.

Clustering and correlation networks within the kinetochore

The fact that proteins in the same complex consistently show very similar H/L ratios across experiments allowed us to perform clustering analysis and map a correlation network of kinetochore proteins (Fig. 9). The clusters seen in Fig. 9 A confirm previous descriptions of CCAN complexes, e.g., CENP-H/-I/-K, CENP-S/-X, CENP-O/-P/-Q/-R/-U, as well as the Mis12, Ndc80, and Ska complexes in the outer kinetochore. They also reveal the existence of two previously undescribed protein cohorts: CENP-N/-L/-T in the inner kinetochore and Rod/Zw10/ Zwilch/Spindly/CENP-E/Mad1 in the outer kinetochore.

The similarity of H/L ratio profiles between pairs of proteins was gauged by Pearson correlation coefficient (Fig. S4). The correlation between every pair of proteins in the chromosomes was summarized as a matrix shown in Fig. 9 B, with a blowup in Fig. S5. A profile of these correlations against the rest of the proteome was used for hierarchical clustering analysis so that proteins with strong correlations are grouped together. One cluster that was enriched for kinetochore proteins was scrutinized in detail (Fig. 9 B). The Mis12 complex, KNL1, and the Ndc80 complex together comprised a single correlation cluster (yellow). This is the KMN network (Cheeseman and Desai, 2008). The Ndc80 complex also formed part of a second cluster together with CENP-N/-L/-T. Interestingly, CENP-N/-L/-T and CENP-H/-I/-K were grouped separately.

In both clustering analysis of H/L ratios and the correlation meta-analysis, CENP-E and Mad1 made a single cluster with the RZZ complex (Fig. 9, A and B). Correlation analysis revealed a link of this RZZ supercohort with CENP-N/-L/-T.
This RZZ cohort also showed a good correlation with the Ndc80 complex, suggesting a significant interaction. On the other hand, the RZZ cohort did not exhibit a high correlation with the Mis12 complex, suggesting that RZZ assembly on chromosomes is linked with Ndc80CENP-T rather than with Ndc80CENP-C.

Discussion

This analysis differs from previous studies of kinetochore assembly in that it exclusively concerns untagged endogenous proteins in situ in native mitotic chromosomes. The results were not constrained by requiring antibodies that may exhibit varying affinities or accessibilities to their target epitopes, tags that might impair protein function in subtle ways, or the need to find experimental conditions where the proteins are soluble.

Dependencies on CENP-A and CENP-C

Neither CENP-A nor CENP-C depends on any other CCAN protein for association with mitotic chromosomes. Depletion of CENP-A and CENP-C was not accompanied by reduction of other proteins to the same extent as themselves, even though CENP-A depletion decreased the chromosomal accumulation of CENP-N/-L/-T and, to a lesser extent, CENP-H/-I/-K/-O/-P/-Q/-R/-U. Thus, CENP-A and CENP-C do not have protein cohorts as defined by this analysis. This is not the case for most kinetochoore components. In the case of the depletion of CENP-H, CENP-I, CENP-K, CENP-M, and CENP-N and loss of CENP-O and CENP-S, one or more other proteins were reduced as much...
as the target protein. Our work supports the view that CENP-A is a marker for the site of kinetochore assembly rather than a structural foundation for the kinetochore (Prendergast et al., 2011).

CENP-N binds to CENP-A (Carroll et al., 2009), and in our experiments, chromosomal association of CENP-N/-L depended on CENP-A. CENP-T/W/S/-X has been reported to bind to DNA independently of CENP-A (Hori et al., 2008a). The unexpected dependency that we observe for CENP-T/-W on CENP-N/-L may favor accumulation of CENP-T/-W near CENP-A nucleosomes. Consistent with this, CENP-N and CENP-T coprecipitated from chromatin extracts as part of the CENP-A associated complex (Foltz et al., 2006) and map at a similar radial position by superresolution microscopy (Suzuki et al., 2014).

CENP-C depletion also had a relatively minor effect on assembly of the inner kinetochore in these experiments. This was surprising in light of the recent study that CENP-C interacts in vitro with a complex of CENP-H/-I/-K/-M (Basilico et al., 2014; Klare et al., 2015) and previous work showing that CENP-C is required for kinetochore localization of CENP-N/-L in yeast (Tanaka et al., 2009; Hinshaw and Harrison, 2013). One possibility is that our depletion of CENP-C (14% remaining on chromosomes) was insufficient to phenocopy the effects seen in previous studies. Alternatively, contributions with other components may also promote the centromere localization of CENP-H/-I/-K/-M in DT40 cells.

CENP-C depletion severely disrupted outer kinetochore assembly. Levels of the entire KMN network were severely reduced on CENP-C–depleted chromosomes. This fits with the view of CENP-C as an essential link between the inner and outer kinetochore (Przewloka et al., 2011; Screpanti et al., 2011).

A network constructed from the correlation matrix based on SILAC ratios for all proteins provides a graphic demonstration of relationships between kinetochore proteins and integrates the diverse experiments reported here (Fig. 10 A). Importantly, no novel proteins clustered among the CCAN members, suggesting that all members of this group of proteins are now known.

The correlation analysis splits “core” CCAN components into the CENP-H/-I/-K/-M complex and CENP-N/-L/-T/-W...
cohort. These are defined not only by their coordinated behavior when constituent kinetochore proteins are deleted but also functionally. CENP-N/-L/-T/-W exhibit a strong interdependency with members of the RZZ cohort (which includes CENP-E, Mad1, and Spindly [see next section]), whereas CENP-H/-I/-K/-M do not.

CENP-O/-P/-Q/-R/-U and CENP-S/-X are distinct complexes that appear to be “peripheral” in the sense that they do not correlate nearly as tightly with the other CCAN components as they do within themselves. Depletion of CENP-O/-P/-Q/-R/-U actually caused an increase in chromosomal levels of CENP-H/-I/-K/-N/-L/-T/S. This was not accompanied by an increase in the KMN and RZZ complexes. These results suggest that CENP-O/-P/-Q/-R/-U may have a role in limiting the size of the inner kinetochore and that the size of the outer kinetochore does not scale directly with the content of CCAN proteins.

The Ndc80 complex as a hub of kinetochore dependencies

Remarkably the Ndc80 complex appears to be a “hub” of the kinetochore dependency network. This complex provides a physical link between kinetochores and dynamic spindle microtubules and has roles in microtubule capture, monitoring the state of microtubule binding, (Alushin et al., 2010, 2012), error correction (Cheeseman et al., 2006; DeLuca et al., 2006), and subsequent transport of chromosomes into daughter cells upon anaphase onset (Tooley and Stukenberg, 2011). Ndc80 is the only complex that shows robust correlations with all other cohorts and complexes diagrammed here (Fig. 10 A). For example, behavior of the CENP-N/-L/-T/-W cohort does not correlate with the Mis12 complex, and CENP-H/-I/-K/-M complex behavior does not correlate with either the Mis12 or RZZ complexes. The Ndc80 complex also shows robust interactions with the Ska complex, which was not included in Fig. 10 A because its interactions as deduced from correlation analysis are surprisingly complex.

We noted from analysis of the variance across all experiments that the Ndc80 complex does not always covary with the other components of the KMN network (the Mis12 complex and Knl1). Indeed, two different types of Ndc80 supercomplex had previously been suggested (Gascoigne et al., 2011; Schleiffer et al., 2012; Nishino et al., 2013). We will refer to these as Ndc80\(^{\text{CENP-C}}\) and Ndc80\(^{\text{CENP-T}}\). The spatial distribution of CENP-C and CENP-T on centromeres is different (Ribeiro et al., 2010), and it was suggested from reconstitution experiments that those supercomplexes might have distinct functions (Hori et al., 2013). In Ndc80\(^{\text{CENP-C}}\), CENP-C binds to the Mis12 complex and Knl1, thereby forming the KMN network. In Ndc80\(^{\text{CENP-T}}\), Spc24 and Spc25 bind CENP-T directly upon phosphorylation of the CENP-T N-terminus (Nishino et al., 2013). The Mis12 complex competes with CENP-T for this binding site. Thus, Ndc80\(^{\text{CENP-T}}\) does not associate with Mis12 and Knl1 and therefore does not form a KMN network.

Our data suggest that Ndc80\(^{\text{CENP-C}}\) and Ndc80\(^{\text{CENP-T}}\) may serve different functions in the kinetochore (Fig. 10 B). Ndc80\(^{\text{CENP-C}}\) exhibits dependency relationships consistent with interactions with the KMN network. It may therefore function primarily in microtubule binding, although a role in the spindle checkpoint has also been suggested (Kim and Yu, 2015). It was previously suggested that the RZZ complex might associate with the KMN network (Varma et al., 2013); however, our dependency analysis suggests a closer functional association between Ndc80\(^{\text{CENP-T}}\) and the expanded RZZ cohort. We do not see dependency interactions between the RZZ cohort and Mis12 components.

The RZZ cohort revealed by our analysis consists of RZZ components plus Spindly, CENP-E, and Mad1. These proteins exhibit such tight correlations with one another that they are likely to form a physical complex in chromosomes. We propose terming this the RZZ--MES (Rod, Zw10, Zwilch, Mad1, CENP-E, Spindly) supercomplex. This complex has not been observed in solution and may only form when its constituents are associated with chromosomes. RZZ is involved in recruiting Mad1/Mad2 to kinetochores (Karess, 2005), and Mad1 in turn recruits CENP-E (Akera et al., 2015). CENP-E is required for conversion to microtubule attachment (Kaspen et al., 2005). MAD1 could cooperate with the RZZ--MES to coordinate end-on microtubule attachments with spindle checkpoint signaling. It will be interesting in future experiments to determine whether Mps1 kinase preferentially associates with Ndc80\(^{\text{CENP-T}}\) (Hiruma et al., 2015; Ji et al., 2015).

Perspectives

This study of the entire mitotic chromosome proteome in a comprehensive set of inner kinetochore mutants has led to the elucidation of a network of protein dependencies throughout the kinetochore. This network divides the CCAN into four
subcomplexes that appear to be relatively independent of one another for localization to mitotic chromosomes (Fig. 10). Perhaps surprisingly, we identified close functional links between CENP-N/-L and CENP-T/-W consistent with them acting as a complex in the kinetochore. Our data also suggest that the RZZ complex may actually be part of a larger RZZ–MES supercomplex that coordinates microtubule attachments with checkpoint signaling in the outer kinetochore. This analysis will hopefully stimulate future biochemical and cell biology experiments to explore the novel relationships identified here.

Materials and methods

Cell culture and medium

The chicken lymphoma B cell line DT40 was grown in RPMI1640 medium supplemented with 10% FBS and 1% penicillin/streptomycin. For SILAC, wild type (clone 18), CENP-C/-H/-M/-N or Mis18α were turned off by treatment with doxycycline for 48 h before harvest of cells. CENP-C/-H/-M/-N or Mis18α were turned off by treatment with doxycycline 12 h before harvest.

Expression of conditional KO cell lines. We refer to such doxycycline-OFF (Fukagawa et al., 2001), CENP-I ON/OFF (Nishihashi et al., 2002), CENP-K ON/OFF (Amano et al., 2009), and CENP-S ON/OFF are conditional KOs (Perpelescu et al., 2015).

For SILAC, lysine and arginine. 100 μg/ml 15C6, 15N2-L-lysine: 2 HCl and 30 μg/ml 15C6, 15N2-L-arginine: HCl were added to the medium with dialyzed FBS. 1 μg/ml doxycycline was used to silence the rescuing cDNA expression of conditional KO cell lines. We refer to such doxycycline-treated cells as GENENAME ON/OFF (e.g., as CENP-C ON/OFF).

Cell lines

Details of the following KO cell lines are described in the original publications: CENP-C ON/OFF (Kwon et al., 2007), CENP-H ON/OFF (Fukagawa et al., 2001), CENP-F ON/OFF (Nishihashi et al., 2002), CENP-K ON/OFF, CENP-M ON/OFF and CENP-O ON/OFF (Okada et al., 2006), CENP-K ON/OFF (Ribeiro et al., 2010), CENP-KG (Hori et al., 2008b), CENP-S ON/OFF (Amano et al., 2009), and HJURP ON/OFF and Mis18 ON/OFF (Perpelescu et al., 2015).

Cell lines designated GENENAME ON/OFF are conditional KOs where all or a portion of an essential gene has been removed and cells are kept alive by a cDNA whose expression can be silenced by addition of doxycycline. Nonessential genes were absolute KOs, designated GENENAME ON/OFF. Expression of exogenous “rescue” cDNAs encoding CENP-C/-H/-M/-N or Mis18α were turned off by treatment with doxycycline for 48 h before harvest of cells. CENP-F ON/OFF and CENP-K ON/OFF cells were incubated with doxycycline for 36 h. HJURP ON/OFF and cells were treated with doxycycline for 84 h.

In all cases, cells were incubated with nocodazole for the last 12 h before harvest.

Preparation of mitotic chromosomes

DT40 cells were incubated with 0.5 μg/ml nocodazole for 12–13 h before harvest. The length of incubation in doxycycline was calibrated such that a minimum mitotic index of >75% was achieved. Longer incubations resulted in increased apoptosis and a decreased mitotic index such that a minimum mitotic index of >75% was achieved. Longer incubations lasted 12 h before harvest.

Mass spectrometry

The DNA content of chromosome isolates was quantified with propidium iodide. Mitotic chromosomes were isolated in SDS-PAGE sample buffer (50 mM Tris, pH 6.8, 2% SDS, 10% glycerol, 0.01% bromophenol blue, and 5% 2-mercaptoethanol). Coomassie-stained gel bands were excised and proteins were digested with trypsin, as previously described (Shi et al., 1999). Briefly, proteins were reduced in 10 mM dithiothreitol (Sigma-Aldrich) for 30 min at 37°C and alkylated in 55 mM iodoacetamide (Sigma-Aldrich) for 20 min at ambient temperature in the dark. They were then digested overnight at 37°C with 12.5 ng/μl trypsin (Pierce). After digestion, samples were acidified with orthophosphoric acid and subjected to strong cation exchange fractionation, using a polySULFOETHYL A column (PolyLC) on an Ultimate 3000 Dionex LC system (Thermo Fisher Scientific). Mobile phase A consisted of 5 mM KH2PO4, 10% acetonitrile at pH 3; mobile phase B was 5 mM KH2PO4, 1 M KCl, and 10% acetonitrile, pH 3. The peptides were fractionated using the following gradient: 0%–60% buffer B in 18 min, then to 70% in 2 min, and then to 0% in 6 min. The flow rate was constant at 200 μl/min. Fractions were collected at 1-min time slices.

For strong cation exchange chromatography, fractions were diluted with an equal volume of 0.1% TFA and spun onto StageTips as described by Rappsilber et al. (2003). Peptides were eluted in 20 μl 80% acetonitrile in 0.1% TFA and concentrated down to 4 μl by vacuum centrifugation (Concentrator 5301; Eppendorf). The peptide sample was then prepared for liquid chromatography tandem mass spectrometry analysis by diluting it to 5 μl by 0.1% TFA. Mass spectrometry analyses were performed either on a Velos LTQ-Orbitrap mass spectrometer (Thermo Fisher Scientific) or on a QExactive Mass spectrometer (Thermo Fisher Scientific), both coupled on-line to Ultimate 3000 RSLCnano Systems (Dionex; Thermo Fisher Scientific).

The analytical column with a self-assembled particle frit (Ishihama et al., 2002) and C18 material (3 μm; ReproSil-Pur C18-AQ; Dr. Maisch GmbH) was packed into a spray emitter (75-μm ID, 8-μm opening, 300-nm length; New Objective) using an air-pressure pump (Proxeon Biosystems). Mobile phase A consisted of water and 0.1% formic acid, and mobile phase B consisted of 80% acetonitrile and 0.1% formic acid. The gradient used was 220 min. In both mass spectrometers, the peptides were loaded onto the column at a flow rate of 0.5 μl/min and eluted at a flow rate of 0.2 μl/min according to the following gradient: 2%–40% buffer B in 180 min, then to 95% in 16 min.

For the LTQ Orbitrap Velos, Fourier transform mass spectrometry spectra were recorded at 60,000 resolution and the 20 most intense peaks with charge ≥2 of the MS scan were selected with an isolation window of 1.0 Da in the ion trap for MS2 (normal scan, wideband activation, filling 5.0E5 ions for MS scan, 1.0E4 ions for MS2, maximum fill time 100 ms, dynamic exclusion for 120 s). For the Q Exactive, FTMS spectra were selected and fragmented by higher-energy collisional dissociation (Ishihama et al., 2007) with normalized collision energy of 25. The maximum ion injection time for the MS and MS2 scans was set to 20 and 60 ms, respectively, and the AGC target was set to 1 E6 for the MS scan and to 5 E4 for the MS2 scan. Dynamic exclusion was set to 60 s.

The MaxQuant software platform (Cox and Mann, 2008) version 1.5.1.2 was used to process the raw files and search was conducted against chicken complete/reference proteome set of UniProt database (released on May 14, 2014) with additional sequences from our in-house database of chicken proteins, using the Andromeda search engine (Cox et al., 2011). The first search peptide tolerance was set to 20 ppm, whereas the main search peptide tolerance was set to 4.5 ppm. Isotope mass tolerance was 2 ppm, and maximum charge was 7. A maximum of two missed cleavages was allowed. Carbamidomethylation of cysteine was set as a fixed modification. Oxidation of methionine and acetylation of the N terminus were set as variable modifications. Peptide and protein identifications were filtered to 1% false discovery rate.
Processing of data
The combined data from 38 experiments contained 2,760 protein IDs. To minimize noise, we removed protein entries that occurred in <25 experiments. 1,667 proteins remained after this filtering process (Table S1). Unprocessed data were used to draw scatter plots and histograms.

In cases where discrepancies in protein H/L ratio were found in the replicates, we inspected the MS1 data manually. Where MS1 peaks suffered from interference because of coeluting signals, the corresponding peptide H/L ratio was removed from the analysis. Centromere proteins affected by this data curation were CENP-M, CENP-R, CENP-U, CENP-W, CENP-X, APC4, and APC5.

Filtering out contaminant proteins
To compare the proteomes of mutant and wild-type chromosomes, it was necessary to differentially label cells with stable isotopes (SILAC) in order to distinguish the chromosomal proteins of mutant and wild-type cells from one another. The metabolic labeling with heavy amino acids used for SILAC requires growth of the cells in medium with dialyzed FBS. Unfortunately, many DT-40 conditional KO cell lines do not grow in medium with dialyzed FBS. We have been unable, despite repeated efforts, to determine the explanation, and we thus grew most mutant cell lines in medium with complete FBS.

To test whether the proteome of isolated mitotic chromosomes changed when cells were cultured in conventional medium or medium with dialyzed FBS, we determined the proteome of wild-type mitotic chromosomes isolated from cells grown under the two sets of conditions. The histogram in Fig. S1 A shows the distribution of log2(H/L ratios) for the 1,667 proteins quantified in this experiment. The distribution was unimodal but skewed to the left. Multipeak analysis identified a second minor peak, comprising ~15% of the total and centered at 2−0.455 ± 0.55 (Fig. S1, A and C).

Cluster analysis with our complete dataset identified three subgroups whose H/L ratios fluctuated in a manner different from the rest (Fig. 1). The first subgroup contained CCAN proteins. The second subgroup is highlighted as purple trees at the top of Fig. 1. They were probably contaminants, many of which were not identified in wild-type mitotic chromosomes. They may be derived from interphase or apoptotic cells. The third group, also highlighted as purple trees, clustered at the bottom of Fig. 1. This group contained proteins that accumulated on chromosomes from cells grown with FBS relative to those grown in dialyzed FBS. The distribution of their H/L ratios coincides with the second minor peak (Fig. S1, B and E). These proteins were almost exclusively identified as cytoplasmic in our previous analysis (Fig. S1 F; Ohta et al., 2010). Because the proteins of the second minor peak were predominantly cytoplasmic, we filtered this set of 259 proteins from our subsequent analyses. This filtration was important, as it decreased random scatter in our data and allowed us to more accurately determine the variance of bona fide chromosomal proteins.

Profile plots
A profile plot is an illustrative summary of changes in the abundance of proteins across experiments (Fig. 4 B). The H/L ratio of a protein in one experiment is plotted along a vertical line, in this case with positive H/L ratios below the x axis so that descending lines reflect decreasing values of the protein in question. Members of the same protein complex typically follow a very similar pattern. The superimposed box plots with whiskers provide a convenient criterion to assess the significance of changes in the abundance or H/L ratio of each protein. The interquartile box contains 50% of all H/L ratios measured in each experiment, with the whiskers encompassing more than 95% of all the total ratios (assuming a normal distribution of values).

Online supplemental material
Fig. S1 shows for several representative experiments the distribution of total proteins and centromere proteins as well as showing the distribution of contaminating proteins that were removed from our statistical analysis. Fig. S2 demonstrates a step-by-step correlation analysis between a pair of experiments with a good correlation (CENP-H^{OFF} vs. CENP-P^{OFF}) and a pair of experiments with a poor correlation (CENP-H^{OFF} vs. CENP-N^{OFF}). Fig. S3 shows profile plots for the CPC, core histones, and the cohesin complexes. Fig. S4 is a step-by-step correlation analysis between the RZZ component Rod and the Mis12 component Nsl1 (poor correlation) and CENP-T (good correlation). Fig. S5 is a blowup of Fig. 9 B, with the positions of several key complexes indicated. Table S1 lists the H/L ratios of all proteins in every experiment. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.201508072/DC1.

Acknowledgments
This work was funded by The Wellcome Trust, of which W.C. Earnshaw is a Principal Research Fellow (grant 107022) and J. Rappsilber is a Senior Research Fellow (grant 103139). The Wellcome Trust Centre for Cell Biology is supported by core grant numbers 077707 and 092076 and instrument grant 091020.

The authors declare no competing financial interests.

Submitted: 18 August 2015
Accepted: 11 November 2015

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Figure S1. Identifying contaminating proteins that were removed from statistical analyses. (A) Histograms show the distribution of log₂(H/L ratio) for all proteins in individual SILAC experiments, with proteins whose levels decreased on chromosomes yielding positive values on the abscissa. Wild-type chromosomes were labeled with heavy amino acids, and KO chromosomes were labeled with light amino acids. (For an as-yet-unknown reason, many of the KO cell lines will not grow in medium with dialyzed serum.) All quantified proteins (gray) are shown for representative wild-type, CENP-I\textsuperscript{OFF}, CENP-H\textsuperscript{OFF}, and CENP-O\textsuperscript{KO} experiments. Bin size is 0.2. The shaded areas span −0.6 to 0.4. (B) As in A, but with “noise” proteins highlighted in red. (C) Histograms after removal of the noise proteins. (D) Behavior of all centromere proteins (blue), with CENP-A and CCAN proteins (red). (E, top) Histogram of log₂(H/L ratio) comparing proteins of chromosomes isolated from wild-type cells grown with dialyzed FBS (heavy amino acid label) versus cells grown with complete FBS (light amino acid label). Red line traces the contour of the histogram. Bin size is 20.2. (E, bottom) Multipeak analysis was applied to the distribution of all quantified proteins. The distribution of log₂H/L was fitted with two Gaussian curves. (F) Classifier II (the ratio of amounts of each protein in cytoplasm versus that in isolated chromosomes; Ohta et al., 2010) was plotted against the log₂(H/L ratio) from the same experiment as E. Virtually all proteins in the small subsidiary peak are known to be cytoplasmic.
Figure S2. Use of correlation analysis to compare results from all experiments. (A) SILAC ratios are plotted for CENP-I \textsuperscript{OFF} chromosomes (abscissa) versus CENP-H \textsuperscript{OFF} chromosomes (ordinate). The behavior of the CCAN proteins (red) shows a high correlation between the two experiments (Pearson coefficient 0.983). (B) SILAC ratios are plotted for CENP-N \textsuperscript{OFF} chromosomes (abscissa) versus CENP-H \textsuperscript{OFF} chromosomes (ordinate). The divergent behavior of some CCAN proteins (red) leads to a poor correlation between the two experiments (Pearson coefficient 0.594). (C) First step in calculating the Pearson correlation coefficient. Each table entry is a diagrammatic plot of the distribution of CCAN proteins exactly as seen in A. It is necessary to blow this up online to see the pattern. The calculated correlation coefficients make up a table of numbers that maps directly onto the diagrammatic table shown in C and are not shown here. Instead, D shows a color-coding of that table. E and F correspond to C and D, but in this case, the plots in E correspond to all centromere proteins shown in blue and red in A. G and H correspond to C and D, but in this case, the plots in G correspond to all proteins shown in A. In the colored matrices shown in D, F, and H red shows a high positive correlation, and green shows a high negative correlation. The matrix (H) is shown in Fig. 2, wt, wild type.
Figure S3. **Coordinated behavior among members of protein complexes.** (A) Profile plot showing the behavior of core histones (H2A, H2B, H3.2, and H4 [red]) in all experiments. (B) Profile plot showing the behavior of chromosomal passenger complex components (Aurora B, INCENP, survivin, and borealin [red]) in all experiments. (C) Profile plot showing the behavior of cohesin subunits (SMC1, SMC3, RAD21, STAG1, and STAG2 [red]) in all experiments. Gray lines show behavior of other proteins.

Figure S4. **Correlation revealed by comparisons in the abundance of two proteins on chromosomes as a result of depletion of CCAN proteins.** (A) Scatterplots showing coordinated behavior of two proteins across all the experiments. Each dot represents respective H/L ratios of a pair of proteins in one experiment. Axes are plotted in a log scale. (B) Pearson correlation coefficients were calculated from the distribution of paired H/L ratios and translated into a coded color. Yellow tiles in Fig. 9 B indicate coordinated behavior of a protein cohort throughout this study.
Figure S5. **Blowup of the correlation matrix shown in Fig. 9 B.**

Provided online as an Excel table is Table S1, which shows the H/L ratio for a protein in each experiment. NaN is a missing value where the protein was not quantified in the given experiment. Proteins marked with a + in the exclusion list column were removed from the analysis.

**Reference**